Temporal variation in loggerhead turtle strandings from Georgia

Final Report to Georgia Department of Natural Resources

Prepared by

Anna L. Bass¹, Shiao-Mei Chow¹, and Brian W. Bowen^{1,2}

¹Department of Fisheries and Aquatic Sciences 7922 NW 71st Street Gainesville, Florida 32653

and

²Archie Carr Center for Sea Turtle Research 223 Bartram Hall University of Florida Gainesville Florida 32611

5 October 1999

INTRODUCTION

Marine turtles utilize different habitats throughout their developmental stages. Knowledge of the origin of animals in different habitats can benefit management by providing information as to which populations may be impacted by human activities. Stranded turtles offer an opportunity to gather information on the populations of turtles that inhabit or travel through different coastal areas. The relative ease of collection of tissue samples also affords researchers the opportunity to test temporal variation in marine turtle cohorts. For example, does the demographic composition of stranded turtles vary seasonally or from year to year?

Loggerhead turtles were previously sampled by stranding network volunteers in 1997 to determine the nesting populations that contribute individuals to Georgian coastal waters. The initial sampling in 1997 provided evidence that at least two nesting populations contribute individuals at detectable levels to the stranded cohort in Georgia (Bass et al., 1998). We also quantified sampling errors in both the mixture (stranded cohort) and stock (nesting populations) samples which affects the precision of contribution estimates from potential source populations.

Samples were collected in 1998 to enhance the sample size and to determine if temporal variation in the stranded cohort could be detected. Investigations in other species of marine turtles have indicated that there is a temporal component to the demographic composition of populations at particular foraging locations (Bass et al., unpubl. data). Because there is some sharing of haplotypes among nesting locations, it is not possible to assign individuals to a given location. The maximum likelihood programs that are currently being used generate the most likely contributions of source populations based on the haplotype frequencies in the stocks and the mixture. The ability of these programs to find the correct answer is based on several assumptions. The most important assumption is that all potential source populations have been characterized and that there are strong shifts in haplotype frequencies in these source populations (Pella and Milner, 1987). Another assumption is that the mixture has been sampled in a manner sufficient to capture the real diversity of haplotypes present in the foraging or stranded population. Investigations into the stock structure of fish have shown that increases in sample size can provide several benefits, such as a reduction in the standard errors of frequency estimates for common haplotypes and an increase in the probability of detecting rare "endemic" haplotypes which in turn increases resolution (Epifanio et al., 1995). Due to the predominance of haplotypes A and B (haplotypes found in the majority of nesting locations used in this analysis) in the 1997 sample and the low frequency of endemic haplotypes, an increase in the sample size of the stranded turtles was desirable to determine if our sampling was a sufficient estimate of the genetic diversity of stranded loggerheads in Georgia.

METHODS

Tissue samples were collected by Sea Turtle Stranding Network volunteers and placed in 15 ml of saturated salt preservation buffer. Samples were then transferred to the University of Florida for analysis. Standard phenol/chloroform DNA isolation protocols were conducted on the tissue samples and a 380 bp fragment of the mitochondrial DNA control region was amplified using primers designed for sea turtles (Allard et al., 1994; Norman et al., 1994). Individual fragments were sequenced and compared to known *Caretta caretta* nesting beach haplotypes. Individuals were then assigned a haplotype based on designations from Encalada et al. (1998) and Bolten et al. (1998).

To test for statistical differences among haplotype frequencies between years, chi-square analyses were performed with the program CHIRXC (Zaykin and Pudovkin, 1993) and probabilities were generated using a Monte Carlo randomization procedure (Roff and Bentzen, 1989).

Maximum likelihood (ML) analysis for mixed stock identification (Grant et al., 1980) was used to estimate the contributions of nesting populations to foraging habitat in Georgia. This method estimates the most likely contributions of source populations based on the haplotype frequencies in the source populations and in the mixed population. The maximum likelihood programs GIRLSEM and UCON were used (Masuda et al., 1991). As a starting point in ML iterations using GIRLSEM, it was assumed that all source populations had an equal probability of contributing (i.e. population size, distance from the foraging location, etc. did not have an impact on the percentage of animals recruiting to a particular area). The conditional maximum likelihood estimates from GIRLSEM were then used as the initial guess to generate a point estimate in the unconditional maximum likelihood program, UCON. Standard errors and 95% confidence intervals of the point estimates were generated from 100 bootstraps of the stock and mixture data sets using GIRLSEM (see Pella et al., 1998).

RESULTS AND DISCUSSION

DNA Analysis of 1998 samples

Of the 124 samples collected, 108 samples produced readable sequence and 16 samples did not work, indicating technical difficulties in the recovery of DNA sequencing information. In the sample of 108 individuals, haplotypes A and B were the most common (Table 1). Six of the haplotypes found during the 1997 sampling period were seen in 1998, but an additional 2 haplotypes (E and J) were found in the 1998 sample. Haplotype E has been observed in the South Florida nesting population while haplotype J has been observed in Mexico. One individual in both the 1997 and 1998 sample sets carried haplotype M (Table 1). In the 1998 set of samples, four individuals which possess haplotype N were found. Haplotypes M and N still remain unknown as to their origin. In addition, four individuals had haplotypes not previously observed at surveyed nesting localities or foraging locations. These were designated GA-1 to GA-4 (see Appendix 1). Although several different haplotypes were found in the second sample represented by the 1998 group, the haplotype frequencies of the two years were not significantly different from each other (X² = 10.20, P = 0.341).

Table 1. Haplotype composition and frequency of stranded loggerhead turtle samples from 1998 and 1997 and locations where these haplotypes have been observed. See Appendix 1 for a detailed listing of haplotypes by turtle or incident number.

| Haplotype | 1998 | 1997 | Location |
|--------------|-------------|-------------|--|
| | | | Nesting ¹ |
| A | 56 | 51 | NWFL, SFL, NEFL-NC |
| В | 33 | 35 | NWFL, SFL, NEFL-NC, Mexico, Greece |
| C | 4 | 6 | NWFL, SFL, Mexico |
| \mathbf{E} | 2 | 0 | SFL |
| G | 2 | 1 | NWFL, SFL |
| J | 2 | 0 | Mexico |
| | (n = 99) | (n = 93) | Foraging ² or Stranded ³ |
| K | 0 | 1 | Madeira ² |
| M | 1 | 1 | Azores ² |
| N | 4 | 6 | Azores ² , Madeira ² , North Atlantic Coast ³ |
| GA-1 | 1 | 0 | Georgia ⁴ |
| GA-2 | 1 | 0 | Georgia ⁴ |
| GA-3 | 1 | 0 | Georgia ⁴ |
| GA-4 | 1 | 0 | Georgia ⁴ |
| | Total = 108 | Total = 101 | |

Abbreviations according to Encalada et al. (1998): NWFL = northwest Florida (panhandle region), SFL = southeast and southwest Florida, NEFL-NC = northeast Florida to North Carolina.

Maximum Likelihood Analysis

Due to the non-significant difference of the haplotype frequencies between the two sampling periods (1997 & 1998), the samples from both years were combined to estimate the contributions of potential source populations (Table 2). This analysis once again provided evidence that SFL contributes the majority of individuals that strand along the Georgia coast. In addition, the nesting population in NWFL was again implicated as a contributor to the stranded turtle cohorts in Georgia.

Table 2. Maximum likelihood estimates of contribution by source populations to stranded loggerhead cohorts from both 1997 and 1998 (n = 192). Estimates were generated using UCON. Standard errors and 95% confidence intervals were generated from 100 bootstraps of both the stock and mixture using GIRLSEM.

| Source population | Contribution (PE) | S.E. | 95% C.I. |
|-------------------|-------------------|--------|---------------|
| NWFL | 0.1999 | 0.1975 | 0.1607-0.2391 |
| SFL | 0.6623 | 0.2028 | 0.6221-0.7025 |
| NEFL-NC | 0.1049 | 0.1269 | 0.0797-0.1301 |
| MEXICO | 0.0327 | 0.0340 | 0.0259-0.0667 |
| BRAZIL | 0 | 0 | |
| GREECE | 0.0000 | 0.0854 | 0.0000-0.0169 |

The ML analysis yielded a notably large estimate (0.1999) from Northwest Florida (Table 2). We believe this result was an overestimate due to in part to the frequency of individuals with haplotype C in the Georgia stranded cohort. After some discussion, we concluded that the estimated contribution from the Florida panhandle (NWFL) was not biologically realistic. Nesting effort in this area may include 100-200 turtles annually (Meylan et al. 1995), as compared to tens of thousands of turtles that nest in southern Florida. Hence the panhandle is an important nesting area, but probably too small to detect with precision in ML analyses. Based on these considerations, we provisionally removed the NWFL rookery from the analysis (Table 3).

Table 3. Maximum likelihood estimates of contribution by source populations to stranded loggerhead cohorts from both 1997 and 1998 (n = 192). The source population NWFL, was removed from the analysis. Estimates were generated using UCON. Standard errors and 95% confidence intervals were generated from 100 bootstraps of both the stock and mixture using GIRLSEM.

| Source Population | Contribution (PE) | S.E. | 95% C.I. |
|-------------------|-------------------|--------|---------------|
| SFL | 0.7274 | 0.1499 | 0.6977-0.7571 |

² Bolten et al. (1998), ³ Rankin-Baransky et al. (submitted), ⁴ This study.

| NEFL-NC | 0.2392 | 0.1098 | 0.2174-0.2610 |
|---------|--------|--------|---------------|
| MEXICO | 0.0334 | 0.0455 | 0.0246-0.0422 |
| BRAZIL | 0.0000 | 0.0000 | |
| GREECE | 0.0000 | 0.0541 | 0.0000-0.0107 |

Removal of NWFL as a potential source population reduces the standard error about the mean for all source populations except Mexico (Table 3). In addition, these estimates appear to be more reflective of population sizes and proximity of nesting populations to the Georgia coastal area. These two factors have been identified as potential determinants of foraging ground composition in green turtles (Bass and Witzell, 2000; Lahanas et al., 1998).

We feel that the estimates of contribution listed in Table 3 are the best estimates for the Georgia stranding cohort.

CONCLUSIONS

Temporal Variation

There is no significant difference among haplotype frequencies from strandings in 1997 (n = 101) and 1998 (n = 108). The main difference between the two sampling years is the presence or absence of low-frequency haplotypes. For example, we observed several rare haplotypes in 1998 that were not detected in 1997, notably haplotypes E and J. Even though we are using a program that attempts to compensate for small sample sizes in some categories (Roff and Bentzen, 1989), the limitations of the Chi-square test may prohibit the detection of subtle differences. We conclude that there are no detectable differences between the 1997 and 1998 stranding cohorts, but this does not eliminate the possibility of temporal variation on longer timescales, or during periods of climatic fluctuation. Sampling from year to year may be too short a time frame to detect temporal variation in loggerheads. Hence it may be useful to reexamine stranded cohorts from Georgia in 5 to 10 years.

Demographic Composition and Comparison to Atlantic Foraging Populations

Out initial analysis yielded an anomalously high contribution from the Florida panhandle. Given the small size of this rookery (Meylan et al. 1995), the estimate of 19% was biologically unrealistic. Subsequent reanalysis without the panhandle population yielded results with lower standard errors. We support the results of this analysis (Table 3) as a basis for management decisions. However, the problem with overestimates from small nesting colonies deserves further attention. One strong possibility is that we have not adequately sampled the genetic variation in regional nesting colonies.

We can state with strong support that a large proportion of the stranded loggerheads are from the south Florida nesting populations. This was apparent in the analysis of samples

from 1997 (77%) and here in the analysis of the combined data set (73%; Table 3). The increased sample size did capture more of the diversity of loggerheads utilizing Georgia waters. The presence of haplotype J also prompted an estimated contribution of 3% from Mexico. Although the standard error is still high and we should be cautious in interpreting this estimate, there do appear to be Mexican loggerheads utilizing US coastal waters.

Investigations of the genetic composition of stranded loggerheads along the northeast coast of the US indicated that 59% were from SFL, 25% from NEFL-NC, and 16% from Mexico (Rankin-Baransky et al., 1999). Notably there was no evidence of a contribution from the NWFL population and the contribution of the SFL population was lower than what we estimate here for the Georgia stranding cohorts. The NC foraging population from Core Sound also exhibits a similar composition: 64% from SFL, 32% from NEFL-NC, and the remaining 4% divided among Mexico and Brazil (Bass et al., 1997). Our estimates for the Georgia stranded cohorts corroborate general conclusions from these other studies. Most strandings along the Atlantic seaboard originate from the SFL nesting population, followed by contributions from NEFL-NC and Yucatan (Mexico). It is likely that other nesting areas contribute at low frequency, but were not detected with current samples and limits of resolution.

Previous research on green turtles, *Chelonia mydas*, has indicated that both nesting population size (Lahanas et al., 1998) and distance from the nesting population to the foraging location (Bass and Witzell, 2000) are correlated with relative contributions to feeding populations. Differences between loggerhead foraging populations may be associated with either of these factors or there may be other unquantified determinants of foraging population composition (Bass and Witzell, 2000).

Management Implications

The estimated composition of the Georgia stranding cohort indicates that several nesting populations may be affected by coastal activities in Georgia. A high proportion of loggerheads derived from the SFL management unit are using Georgian waters as a migratory pathway or resident feeding habitat. In addition about a quarter of the stranded cohort are derived from the NEFL-NC management unit with the remaining proportion composed of animals derived from the Mexican nesting locality. Other diminutive or unsurveyed populations in the Atlantic (such as those in Northwest Florida, the Bahamas, or Cuba) may be affected as well.

What are the management implications for the Georgia nesting population? In the ML analysis, the Georgia nesting population is combined with those in South Carolina, North Carolina, and northeastern Florida (Amelia Island), because we lack the resolution to distinguish these nesting areas with current mtDNA assays. (Forthcoming microsatellite surveys may alleviate this problem.) Hence we cannot give an exact contribution of the Georgia nesting population to the Georgia strandings. However, it is reasonable to assume that Georgia contributes to the 21-26% estimate from NEFL-NC at a level proportional to the size of this nesting population.

ACKNOWLEDGEMENTS

Logistic support was provided by the Archie Carr Center for Sea Turtle Research, and the Department of Fisheries and Aquatic Sciences at the University of Florida. Additional financial support was provided by the Turner Foundation, National Marine Fisheries Service, Ocean Fund, and National Science Foundation. Thanks to the DNA Sequencing Core at the University of Florida. Special thanks to Jerome Pella and Michele Masuda of the Alaska Fisheries Science Center for assistance with data analysis.

LITERATURE CITED

- Allard, M.W., M.M. Miyamoto, K.A. Bjorndal, A.B. Bolten, and B.W. Bowen. 1994. Support for natal homing in green turtles from mitochondrial DNA sequences. Copeia 1994:34-41
- Bass, A.L. and W.N. Witzell. 2000. Demographic composition of immature green turtles (*Chelonia mydas*) from the east central Florida coast: evidence from mtDNA markers. Herpetologica. *In Press*.
- Bass, A.L., S.-M. Chow, and B.W. Bowen. 1998. Nesting cohort affiliation of stranded *Caretta caretta* recovered in Georgia. Final Report to Georgia Department of Natural Resources.
- Bass, A.L., C.J. Lagueux, and B.W. Bowen. 1998. Origin of green turtles, *Chelonia mydas*, at "sleeping rocks" off the northeast coast of Nicaragua. Copeia 4:1064-1069.
- Bass, A.L., S.P. Epperly, J. Braun, and D.W. Owens. 1997. Natal origin and sex ratios of foraging sea turtles in the Pamlico-Albemarle estuarine complex. Poster Presented at 1997 Sea Turtle Symposium, Orlando, Fl.
- Bolten, A.B., K.A. Bjorndal, H.R. Martins, T. Dellinger, M.J. Biscoito, S.E. Encalada, and B.W. Bowen. 1998. Transatlantic developmental migrations of loggerhead sea turtles demonstrated by mtDNA sequence analysis. Ecological Applications, 8(1):107.
- Bowen, B.W. and W.N. Witzell. 1996. Sea turtle conservation genetics. Pp. 1-7. *In* B. W. Bowen and W. N. Witzell (Eds.), Proceedings of the International Symposium on Sea Turtle Conservation Genetics. NOAA Technical Memorandum. NMFS-SEFSC-396. Miami, Florida. U.S.A.
- Bowen, B.W., F.A. Abreu-Grobois, G.H. Balazs, N. Kamezaki, C.J. Limpus, and R.J. Ferl. 1995. Trans-Pacific migrations of the loggerhead turtle (Caretta caretta) demonstrated with mitochondrial DNA markers. Proceedings of the National Academy of Science USA 92:3731-3734.
- Broderick, D., C. Moritz, J.D. Miller, M. Guinea, R.J. Prince, and C.J. Limpus. 1994. Genetic studies of the hawksbill turtle *Eretmochelys imbricata*: evidence for multiple stocks in Australian waters. Pacific Conservation Biology 1:123-131.
- Epifanio, J.M., P.E. Smouse, C.J. Kobak, and B.L. Brown. 1995. Mitochondrial DNA divergence among populations of American shad (*Alosa sapidissima*): how much

- variation is enough for mixed-stock analysis? Canadian Journal of Fisheries and Aquatic Sciences 52:1688-1702.
- Encalada, S.E., K.A. Bjorndal, A.B. Bolten, J.C. Zurita, B. Schroeder, E. Possardt, C.J. Sears, B.W. Bowen. 1998. Population structure of loggerhead turtle (*Caretta caretta*) nesting colonies in the Atlantic and Mediterranean as inferred from mitochondrial DNA control region sequences. Marine Biology 130:567-575.
- Grant, W.S., G.B. Milner, P. Krasnowski, and F.M. Utter. 1980. Use of biochemical genetic variants for identification of sockeye salmon (*Oncorrhynchus nerka*) stocks in Cook Inlet, Alaska. Canadian Journal of Fisheries and Aquatic Sciences 37:1236-1247.
- Lahanas, P.N., K.A. Bjorndal, A.B. Bolten, S.E. Encalada, M.M. Miyamoto, R.A. Valverde, and B.W. Bowen. 1998. Genetic composition of a green turtle feeding ground population: Evidence for multiple origins. Marine Biology 130:345-352.
- Laurent, L., P. Casale, M.N. Bradai, B.J. Godley, G. Gerosa, A.C. Broderick, W. Schroth,
 B. Schierwater, A.M. Levy, D. Freggi, E.M. Abd El-Mawla, D.A. Hadoud, H.E.
 Gomati, M. Domingo, M. Hadjichristophosou, L. Kornaraky, F. Demirayak, and
 C.H. Gautier. 1998. Molecular resolution of marine turtle stock composition in
 fishery bycatch: a case study in the Mediterranean. Molecular Ecology 7:1529-1542.
- Masuda, M., S. Nelson, and J. Pella. 1991. User manual for GIRLSEM, GIRLSYM, AND CONSQRT. USA-DOC-NOAA-NMFS. Programs and user manual available from US-Canada Salmon Program, 11305 Glacier Hwy., Juneau, Alaska 99801 USA.
- Meylan, A., B. Schroeder, and A. Moser. 1995. Sea turtle nesting activity in the state of Florida 1979-1992. Florida Marine Research Publication 52. Florida Marine Research Institute, St. Petersberg, FL.
- Norman, J.A., C. Moritz, C.J. Limpus. 1994. Mitochondrial DNA control region polymorphisms: Genetic markers for ecological studies of marine turtles. Molecular Ecology 3:363-373.
- Pella, J.J. and G.B. Milner. 1987. Use of genetic marks in stock composition analysis. Pp. 247-276 *in* N. Ryman and F. Utter, eds. Population genetics and fisheries management. Univ. of Washington Press, Seattle.
- Pella, J., M. Masuda, C.G. Guthrie, C. Kondzela, A. Gharrett, A. Moles, G. Winans. 1998. Stock composition of some sockeye salmon, *Oncorhynchus nerka*, catches in southeast Alaska, based on incidence of allozyme variants, freshwater ages, and a brain-tissue parasite. U.S. Depart. of Commerce, NOAA Technical Report NMFS 132, 23 p.
- Rankin-Baransky, K., C.J. Williams, A.L. Bass, B.W. Bowen, and J.R. Spotila. 1999. Origin of loggerhead turtle strandings in the northwest Atlantic as determined by mtDNA analysis. *Submitted*.
- Roff, D.A. and P. Bentzen. 1989. The statistical analysis of mitochondrial DNA polymorphisms: Chi-square and the problem of small samples. Molecular Biology and Evolution 6:539-545.
- Zaykin, D.V. and A.I. Pudovkin. 1993. Two programs to estimate significance of X2 values using pseudoprobability tests. Journal of Heredity 84:152.

| Appendix 1 Lab# | . Distribution of haplotypes Stranding ID # | s in stranded individuals Location | Haplotype |
|--------------------|--|-------------------------------------|-----------|
| | SAMPLES FROM 1997 | | |
| 915 | CR-97060601-CC-CI | Georgia; Cumberland Island | В |
| 916 | CR-97053003-CC-CI | Georgia; Cumberland Island | N |
| 917 | CR-97070101-CC-CI | Georgia; Cumberland Island | В |
| 918 | CR-97062601-CC-CI | Georgia; Cumberland Island | N |
| 919 | CR-97072102-CC-CI | Georgia; Cumberland Island | A |
| 920 | CR-97081001-CC-CI | Georgia; Cumberland Island | A |
| 921 | CR-97072501-CC-CI | Georgia; Cumberland Island | В |
| 922 | CR-97060503-CC-CI | Georgia; Cumberland Island | В |
| 923 | CR-97050901-CC-CI | Georgia; Cumberland Island | В |
| 924 | CR-97053001-CC-CI | Georgia; Cumberland Island | В |
| 925 | CR-97062702-CC-CI | Georgia; Cumberland Island | A |
| 926 | CR-97081201-CC-CI | Georgia; Cumberland Island | A |
| 927 | CR-97081301-CC-CI | Georgia; Cumberland Island | A |
| 928 | CR-97091401-CC-CI | Georgia; Cumberland Island | A |
| 929 | CR-97060604-CC-CI | Georgia; Cumberland Island | A |
| 930 | CR-97051701-CC-CI | Georgia; Cumberland Island | В |
| 931 | CR-97060402-CC-CI | Georgia; Cumberland Island | A |
| 932 | CR-97060401-CC-CI | Georgia; Cumberland Island | В |
| 933 | CR-97092202-CC-CI | Georgia; Cumberland Island | A |
| 934 | CR-97092401-CC-CI | Georgia; Cumberland Island | В |
| 935 | CR-97100301-CC-CI | Georgia; Cumberland Island | G |
| 936 | CR-97052501-CC-CI | Georgia; Cumberland Island | В |
| 937 | CR-97052902-CC-CI | Georgia; Cumberland Island | В |
| 938 | CR-97062401-CC-CI | Georgia; Cumberland Island | В |
| 939 | CR-97062801-CC-CI | Georgia; Cumberland Island | В |
| 940 | CR-97072301-CC-CI | Georgia; Cumberland Island | В |
| 941 | CR-97060502-CC-CI | Georgia; Cumberland Island | A |
| 942 | CR-97072101-CC-CI | Georgia; Cumberland Island | A |
| 943 | CR-97051703-CC-CI | Georgia; Cumberland Island | В |
| 944 | CR-97062603-CC-CI | Georgia; Cumberland Island | В |
| 945 | CR-97042604-CC-CI | Georgia; Cumberland Island | C |
| 946 | CR-97062703-CC-CI | Georgia; Cumberland Island | A |
| 947 | CR-97061901-CC-CI | Georgia; Cumberland Island | A |
| 948 | CR-97072201-CC-CI | Georgia; Cumberland Island | A |
| 949 | CR-97062402-CC-CI | Georgia; Cumberland Island | A |
| 950 | CR-97052502-CC-CI | Georgia; Cumberland Island | В |
| 951 | CR-97060501-CC-CI | Georgia; Cumberland Island | A |
| 952 | CR-97071301-CC-CI | Georgia; Cumberland Island | В |
| 953 | CR-97071601-CC-CI | Georgia; Cumberland Island | A |
| 954 | CR-97092101-CC-CI | Georgia; Cumberland Island | A |
| 955 | CR-97061902-CC-CI | Georgia; Cumberland Island | В |
| 956 | CR-97063003-CC-CI | Georgia; Cumberland Island | В |
| 957 | CR-97052303-CC-CI | Georgia; Cumberland Island | Ā |
| 958 | CR-97062501-CC-CI | Georgia; Cumberland Island | C |
| 959 | CR-97052801-CC-CI | Georgia; Cumberland Island | A |
| 960 | CR-97063002-CC-CI | Georgia; Cumberland Island | Α |
| | | 2 / | |

| Lab# | Stranding ID # | Location | Haplotype |
|------|-------------------|----------------------------|-----------|
| 961 | CR-97050101-CC-CI | Georgia; Cumberland Island | A |
| 962 | CR-97052301-CC-CI | Georgia; Cumberland Island | A |
| 963 | CR-97060603-CC-CI | Georgia; Cumberland Island | В |
| 964 | CR-97100102-CC-CI | Georgia; Cumberland Island | A |
| 965 | CR-97092201-CC-CI | Georgia; Cumberland Island | A |
| 966 | CR-97052503-CC-CI | Georgia; Cumberland Island | A |
| 967 | CR-97051301-CC-CI | Georgia; Cumberland Island | A |
| 968 | CR-97070102-CC-CI | Georgia; Cumberland Island | A |
| 969 | CR-97062606-CC-CI | Georgia; Cumberland Island | M |
| 970 | CR-97082401-CC-CI | Georgia; Cumberland Island | A |
| 971 | CR-97051702-CC-CI | Georgia; Cumberland Island | A |
| 972 | CR-97052302-CC-CI | Georgia; Cumberland Island | A |
| 973 | CR-97053002-CC-CI | Georgia; Cumberland Island | A |
| 974 | CR-97062701-CC-CI | Georgia; Cumberland Island | N |
| 975 | CR-97062605-CC-CI | Georgia; Cumberland Island | В |
| 976 | CR-97081801-CC-CI | Georgia; Cumberland Island | В |
| 977 | GA-97052201 | Georgia; JEK | N |
| 978 | GA-97071403 | Georgia; JEK | В |
| 979 | GA-97092302 | Georgia; JEK | В |
| 980 | GA-97092301 | Georgia; JEK | В |
| 981 | GA-97071401 | Georgia; JEK | В |
| 982 | GA-97060501 | Georgia; JEK | A |
| 983 | GA-97080802 | Georgia; JEK | NW |
| 984 | GA-97081201 | Georgia; JEK | A |
| 985 | GA-97061701 | Georgia; JEK | NW |
| 986 | GA-97071201 | Georgia; JEK | NW |
| 987 | GA-97071402 | Georgia; JEK | N |
| 988 | GA-97051501 | Georgia; JEK | NW |
| 989 | GA-97080801 | Georgia; JEK | В |
| 990 | GA-97052401 | Georgia; SAP | С |
| 991 | GA-970701001 | Georgia; SAP | В |
| 992 | GA-97090901 | Georgia; SAP | A |
| 993 | GA-970701601 | Georgia; SAP | A |
| 994 | GA-97052301 | Georgia; SAP | В |
| 995 | GA-97062401 | Georgia; Sea Island | A |
| 996 | GA-97070302 | Georgia; Sea Island | A |
| 997 | GA-97070601 | Georgia; Sea Island | A |
| 998 | GA-97042401 | Georgia; Sea Island | В |
| 999 | GA-97063001 | Georgia; Sea Island | A |
| 1000 | GA-97072301 | Georgia; Sea Island | A |
| 1001 | GA-97062901 | Georgia; Sea Island | В |
| 1002 | GA-97070301 | Georgia; Sea Island | K |
| 1003 | GA-97071301 | Georgia; St. Simons Island | В |
| 1004 | GA-97050801 | Georgia; St. Simons Island | A |
| 1005 | GA-97090101 | Georgia; St. Simons Island | C |
| 1006 | GA-97062601 | Georgia; St. Simons Island | A |

| Lab# | Stranding ID # | Location | Haplotype |
|------|--------------------------|---------------------------|-----------|
| 1007 | GA-97050801 | Georgia ** | NW |
| 1008 | GA-97050601 | Georgia ** | NW |
| 1009 | GA-97070201 | Georgia ** | C |
| 1010 | GA-97052501 | Georgia ** | A |
| 1011 | GA-97052001 | Georgia; OSS | NW |
| 1012 | GA-97062301 | Georgia; OSS | N |
| 1013 | GA-97052401 | Georgia; OSS | В |
| 1014 | GA-97070201 | Georgia; OSS | A |
| 1015 | | Georgia; SCI | NW |
| 1016 | GA-9708071 | Georgia; SCI | NW |
| 1017 | GA-9706131 | Georgia; SCI | C |
| 1018 | GA-97052601 | Georgia; SCI | A |
| 1019 | GA-97051401 | Georgia; LSS | A |
| 1020 | GA-9706091 | Georgia; LSS | A |
| 1021 | GA-97070901 | Georgia; LSS | A |
| 1022 | GA-97082001 | Georgia; LSS | В |
| 1023 | GA-97060601 | Georgia; LSS | NW |
| 1024 | GA-97070601 | Georgia; LSS | A |
| 1025 | GA-97052801 | Georgia; LSS | A |
| | SAMPLES FROM 1998 | | |
| 1863 | GA98091501 | GA; Tybee Isl | В |
| 1864 | GA98070501 | GA; Cumberland Isl | J |
| 1865 | GA98071301 | GA; Tybee Isl | В |
| 1866 | GA98082001 | GA; Tybee Isl | A |
| 1867 | GA98101101 | GA; Little Cumberland Isl | NW |
| 1868 | GA98060301 | GA; Little Cumberland Isl | В |
| 1869 | GA98062501 | GA; Sapelo Isl | A |
| 1870 | GA98070301 | GA; Sapelo Isl | J |
| 1871 | GA98063001 | GA; Sapelo Isl | A |
| 1872 | GA98072201 | Ga; Saint Simons Isl | В |
| 1873 | GA98062801 | Ga; Saint Simons Isl | В |
| 1874 | GA98091101 | Ga; Saint Simons Isl | В |
| 1875 | GA98070101 | GA; Sea Isl | GA-1 |
| 1876 | GA98061401 | GA; Sea Isl | A |
| 1877 | GA98051902 | GA; Saint Catherine's Isl | В |
| 1878 | GA98061101 | GA; Sea Isl | C |
| 1879 | GA98062201 | GA; Sea Isl | A |
| 1880 | GA98081001 | GA; Sea Isl | В |
| 1881 | GA98050601 | GA; Jekyll Isl | В |
| 1882 | GA98050101 | GA; Jekyll Isl | В |
| 1883 | GA98080602 | GA; Jekyll Isl | GA-2 |
| 1884 | GA98092401 | GA: Tybee Isl | В |
| 1885 | GA98052401 | GA; Jekyll Isl | В |
| 1886 | GA98060601 | GA; Tybee Isl | В |
| 1887 | GA98062702 | GA; Jekyll Isl | A |
| 1888 | GA98062701 | GA; Jekyll Isl | A |

| Lab# | Stranding ID # | Location | Haplotype |
|------|----------------|---------------------------|-----------|
| 1889 | GA98052901 | GA; Jekyll Isl | В |
| 1890 | GA98052902 | GA; Jekyll Isl | A |
| 1891 | GA98040601 | GA; Jekyll Isl | В |
| 1892 | GA98090901 | GA; Jekyll Isl | A |
| 1893 | GA98051101 | GA; Jekyll Isl | A |
| 1894 | GA98053001 | GA; Jekyll Isl | A |
| 1895 | GA98051401 | GA; Jekyll Isl | A |
| 1896 | GA98062501 | GA; Saint Catherine's Isl | A |
| 1897 | GA98052901 | GA; Ossabaw Isl | NW |
| 1898 | GA98071801 | GA; Ossabaw Isl | В |
| 1899 | GA98051901 | GA; Ossabaw Isl | NW |
| 1900 | GA98082001 | GA; Ossabaw Isl | GA-3 |
| 1901 | GA98070901 | GA; Ossabaw Isl | NW |
| 1902 | GA98060901 | GA; Ossabaw Isl | A |
| 1903 | GA98052401 | GA; Ossabaw Isl | В |
| 1904 | GA98060801 | GA; Ossabaw Isl | A |
| 1905 | GA98063001 | GA; Ossabaw Isl | NW |
| 1906 | GA98083102 | GA; Ossabaw Isl | NW |
| 1907 | GA98092701 | GA; Cumberland Isl | В |
| 1908 | GA98080401 | GA; Cumberland Isl | A |
| 1909 | GA98092401 | GA; Cumberland Isl | A |
| 1910 | GA98061101 | GA; Cumberland Isl | N |
| 1911 | GA98051901 | GA; Cumberland Isl | M |
| 1912 | GA98070901 | GA; Saint Catherine's Isl | A |
| 1913 | GA98062801 | GA; Cumberland Isl | A |
| 1914 | GA98072102 | GA; Cumberland Isl | N |
| 1915 | GA98062304 | GA; Cumberland Isl | A |
| 1916 | GA98080402 | GA; Cumberland Isl | A |
| 1917 | GA98070601 | GA; Cumberland Isl | G |
| 1918 | GA98091601 | GA; Cumberland Isl | A |
| 1919 | GA98060903 | GA; Cumberland Isl | В |
| 1920 | GA98070602 | GA; Cumberland Isl | A |
| 1921 | GA98061801 | GA; Cumberland Isl | A |
| 1922 | GA98071301 | GA; Cumberland Isl | В |
| 1923 | GA98071301 | GA; Saint Catherine's Isl | A |
| 1924 | GA98072101 | GA; Cumberland Isl | A |
| 1925 | GA98072301 | GA; Saint Catherine's Isl | A |
| 1926 | GA98071001 | GA; Cumberland Isl | A |
| 1927 | GA98070604 | GA; Cumberland Isl | A |
| 1928 | GA98111203 | GA; Cumberland Isl | В |
| 1929 | GA98062301 | GA; Cumberland Isl | A |
| 1930 | GA98071201 | GA; Cumberland Isl | Α |
| 1931 | GA98060102 | GA; Cumberland Isl | A |
| 1932 | GA98072301 | GA; Cumberland Isl | A |
| 1933 | GA98061201 | GA; Cumberland Isl | В |
| 1934 | GA98062101 | GA; Cumberland Isl | NW |

| Lab# | Stranding ID # | Location | Haplotype |
|------|----------------|---------------------------|-----------|
| 1935 | GA98052001 | GA; Cumberland Isl | A |
| 1936 | GA98110901 | GA; Cumberland Isl | A |
| 1937 | GA98070801 | GA; Cumberland Isl | A |
| 1938 | GA98042901 | GA; Cumberland Isl | В |
| 1939 | GA98061001 | GA; Cumberland Isl | NW |
| 1940 | GA98062203 | GA; Cumberland Isl | A |
| 1941 | GA98100101 | GA; Cumberland Isl | E |
| 1942 | GA98091101 | GA; Cumberland Isl | C |
| 1943 | GA98062502 | GA; Cumberland Isl | A |
| 1944 | GA98062305 | GA; Cumberland Isl | A |
| 1945 | GA98060901 | GA; Cumberland Isl | N |
| 1946 | GA98053101 | GA; Cumberland Isl | G |
| 1947 | GA98043002 | GA; Cumberland Isl | NW |
| 1948 | GA98082101 | GA; Cumberland Isl | A |
| 1949 | GA98062303 | GA; Cumberland Isl | В |
| 1950 | GA98050701 | GA; Cumberland Isl | A |
| 1951 | GA98092402 | GA; Cumberland Isl | GA-4 |
| 1952 | GA98062204 | GA; Cumberland Isl | A |
| 1953 | GA98062802 | GA; Cumberland Isl | A |
| 1954 | GA98031601 | GA; Cumberland Isl | В |
| 1955 | GA98060905 | GA; Cumberland Isl | E |
| 1956 | GA98053102 | GA; Cumberland Isl | A |
| 1957 | GA98060904 | GA; Cumberland Isl | A |
| 1958 | GA98072202 | GA; Cumberland Isl | В |
| 1959 | GA98062701 | GA; Cumberland Isl | C |
| 1960 | GA98052101 | GA; Cumberland Isl | В |
| 1961 | GA98060902 | GA; Cumberland Isl | A |
| 1962 | GA98051402 | GA; Cumberland Isl | A |
| 1963 | GA98062302 | GA; Cumberland Isl | В |
| 1964 | GA98041601 | GA; Cumberland Isl | NW |
| 1965 | GA98040202 | GA; Cumberland Isl | C |
| 1966 | GA98053001 | GA; Cumberland Isl | A |
| 1967 | GA98062202 | GA; Cumberland Isl | A |
| 1968 | GA98043001 | GA; Cumberland Isl | NW |
| 1969 | GA98073001 | GA; Cumberland Isl | A |
| 1970 | GA98073002 | GA; Cumberland Isl | A |
| 1971 | GA98060104 | GA; Cumberland Isl | В |
| 1972 | GA98051403 | GA; Cumberland Isl | В |
| 1973 | GA98082001 | GA; Jekyll Isl | A |
| 1974 | GA98072401 | GA; Ossabaw Isl | A |
| 1975 | GA98051903 | GA; Saint Catherine's Isl | В |
| 1976 | GA97052401 | GA; Tybee Isl | A |
| 1977 | GA98060101 | GA; Sant Catherine's Isl | NW |
| 1978 | GA98072302 | GA; Blackbeard Isl | В |
| 1979 | GA98101001 | GA; Blackbeard Isl | NW |
| 1980 | GA98053002 | GA; Blackbeard Isl | A |

| Lab # | Stranding ID # | Location | Haplotype |
|-------|----------------|--------------------|-----------|
| 1981 | GA98053001 | GA; Blackbeard Isl | В |
| 1982 | GA98081701 | GA; Blackbeard Isl | N |
| 1983 | GA98072301 | GA; Blackbeard Isl | A |
| 1984 | GA98053101 | GA; Blackbeard Isl | NW |
| 1985 | GA98062601 | GA; Blackbeard Isl | NW |
| 1986 | GA98060101 | GA; Wassaw Isl | NW |

NW= Not Working